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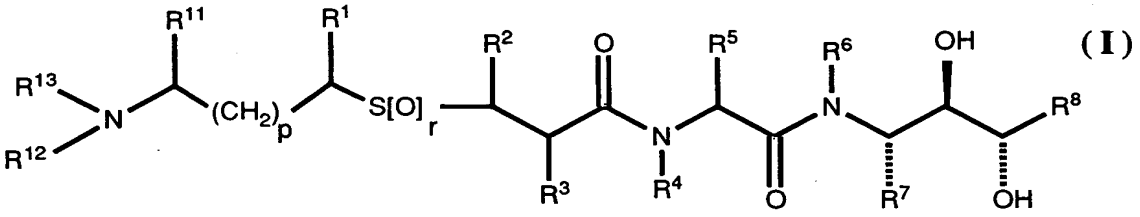
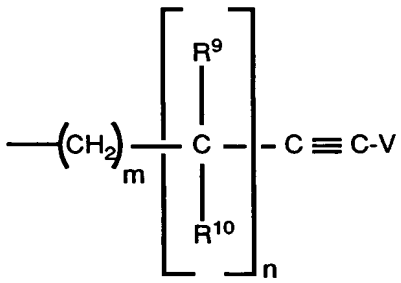
What Is Claimed Is:

1. A compound of Formula I:



(I)

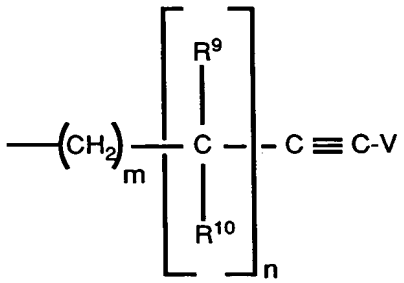
wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl; wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is a group selected from hydrido, cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of R^4 and R^6 is a group independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R^9 and R^{10} is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R^8 is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R^{12} and R^{13} is a group independently selected from hydrido, alkyl, cycloalkyl,

cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R^1 through R^{13} groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

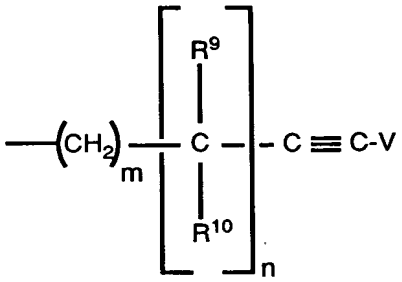
2. Compound of Claim 1 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, iso-butyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'-diethylaminomethyl, N,N'diethylaminoethyl and phenyl; wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R^9 and R^{10} is independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R^8 is selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl

and haloalkenyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkanoyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

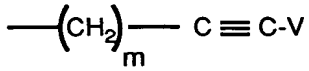
3. Compound of Claim 2 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected from zero through three, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one through three; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl,

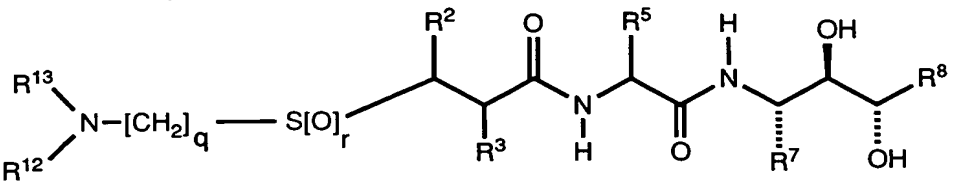
tert-butyl, cyclopropyl, cyclopropylmethyl, cyclopropylethyl, propylcarbonyl, ethylcarbonyl, methylcarbonyl, phenyl, benzyl, phenylethyl, monochlorophenyl, dichlorophenyl, monofluorophenyl, difluorophenyl, monochlorophenylmethyl, monochlorophenylethyl, dichlorophenylmethyl, dichlorophenylethyl, naphthyl, monofluoronaphthyl, monochloronaphthyl, naphthylmethyl, naphthylethyl, fluoronapthylmethyl and chloronaphthylethyl; or a pharmaceutically-acceptable salt thereof.

4. Compound of Claim 3 wherein each of R^1 and R^{11} is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from



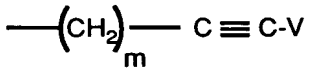
wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R^{12} and R^{13} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

5. Compound of Claim 4 of Formula II



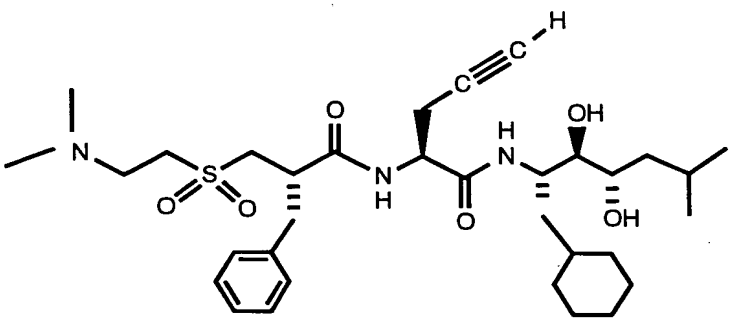
(II)

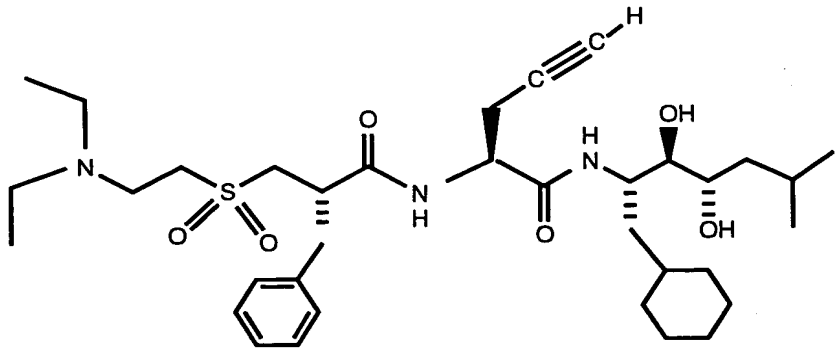
wherein r is zero or two; wherein q is two or three; wherein R^2 is selected from hydrido, methyl, ethyl and phenyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from

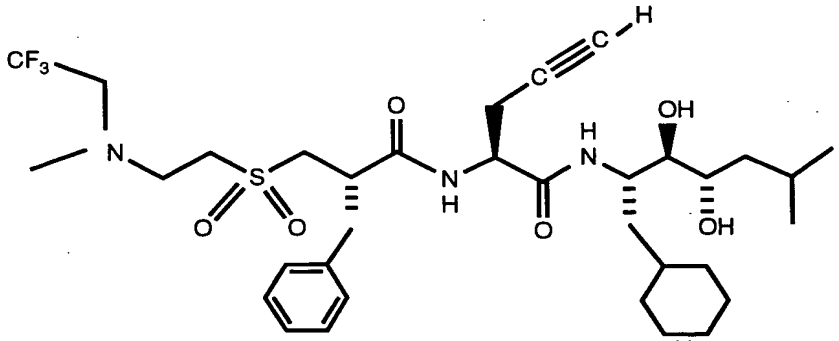


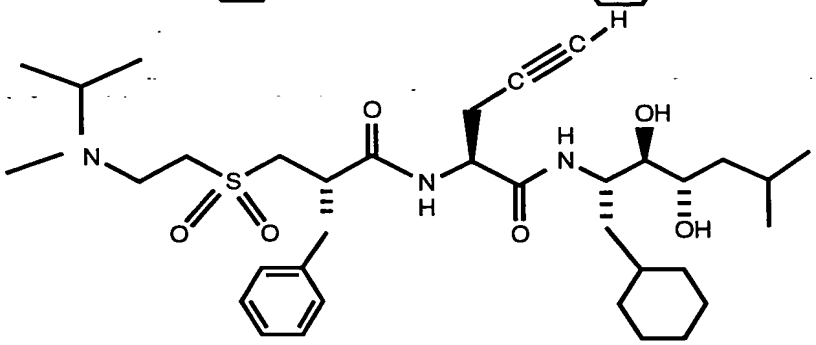
wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R^{12} and R^{13} is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

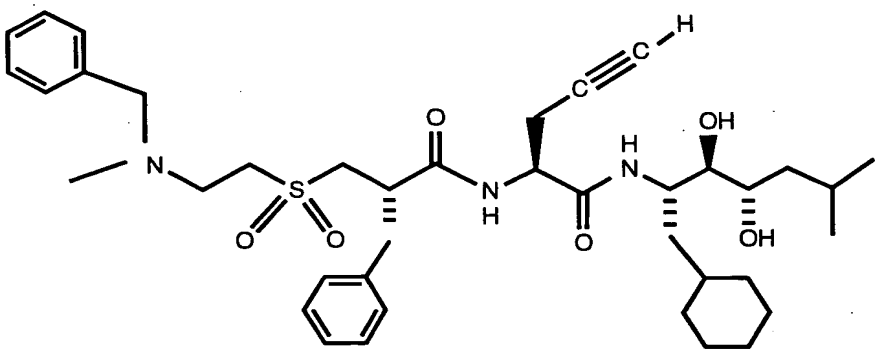
6. Compound of Claim 5 selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

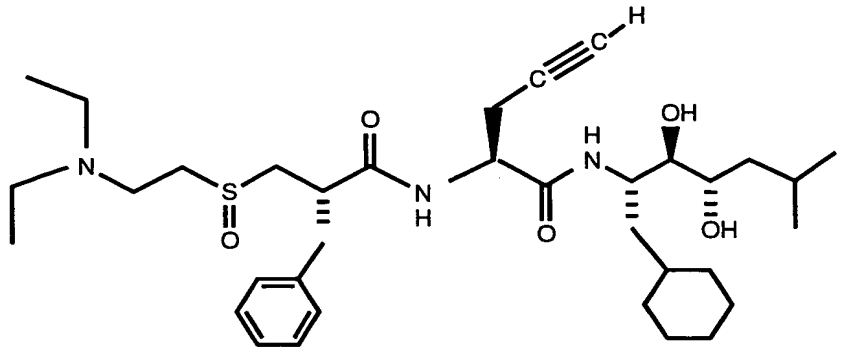


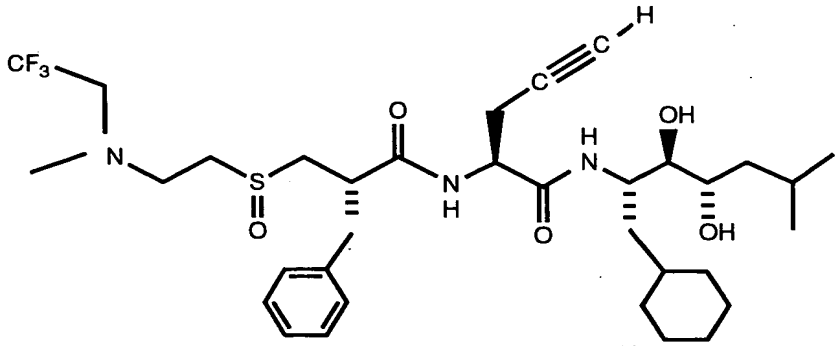


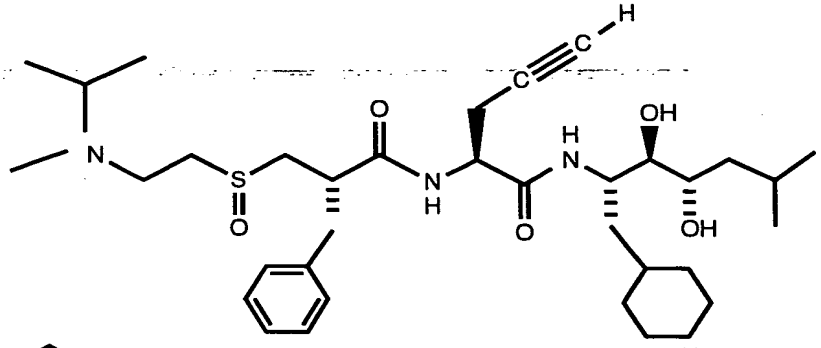


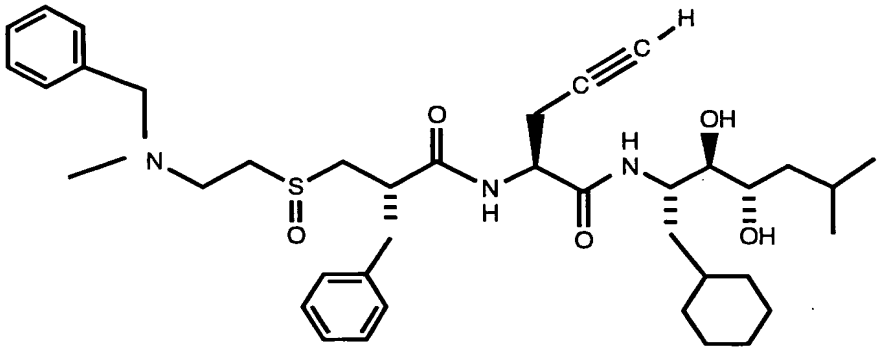


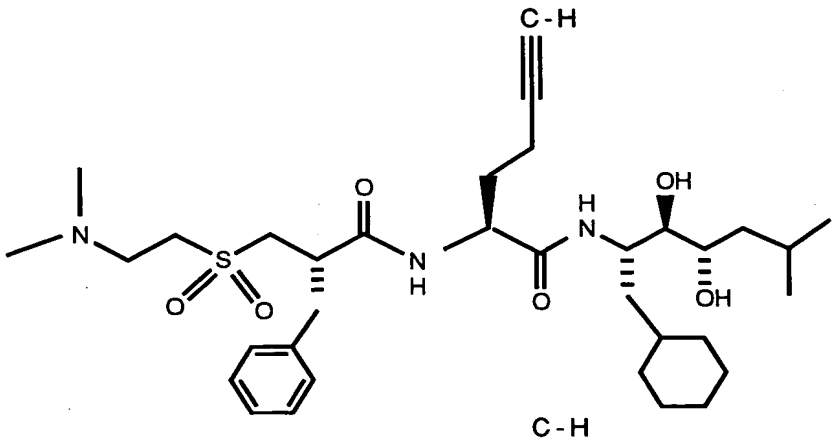


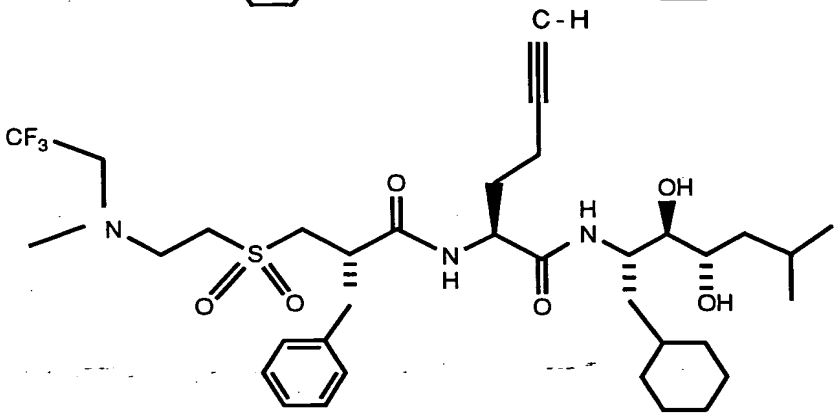


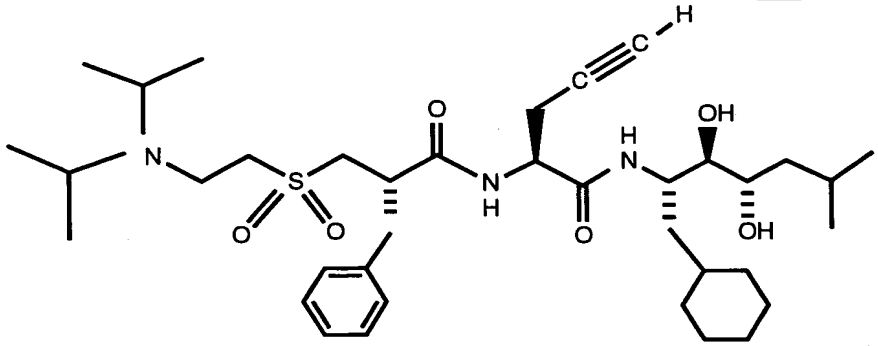




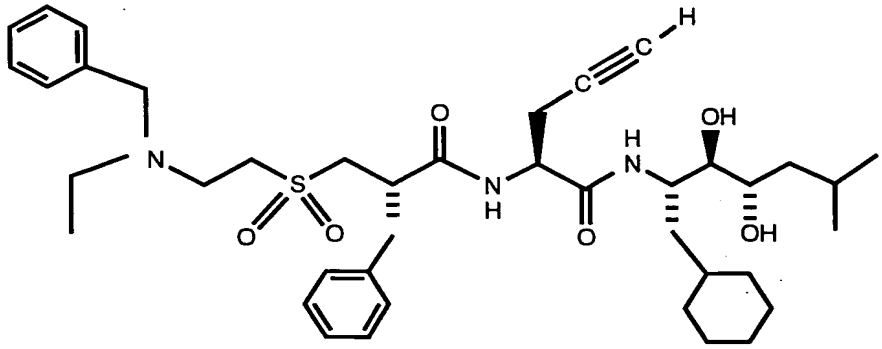








and



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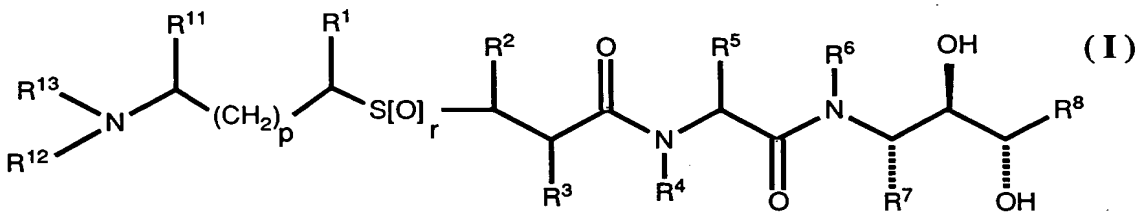
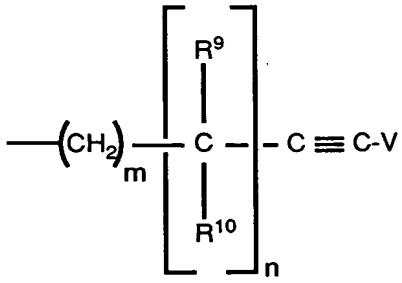
7. Compound of Claim 6 which is N-[1R*-[[[1S,1R*-(cyclohexylmethyl)-2S*,3R*-dihydroxy-5-

methylhexyl]amino]carbonyl]-3-butynyl]-αR*-[[[2-(dimethylamino)ethyl]sulfonyl]methyl]benzenepropanamide or a pharmaceutically-acceptalble salt thereof.

8. A pharmaceutical composition comprising a therapeutically-effective amount of a renin-inhibiting compound and a pharmaceutically-acceptable carrier or diluent, said renin-inhibiting compound selected from a family of compounds of Formula I:



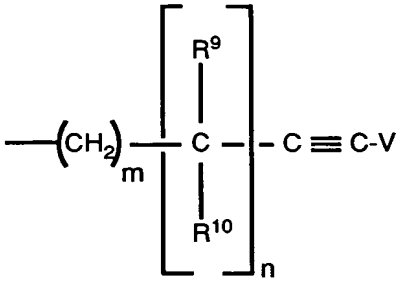
wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl; wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is a group selected from hydrido, cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of R^4 and R^6 is a group independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R^9 and R^{10} is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number

selected from zero through three; wherein R^7 is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R^8 is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R^{12} and R^{13} is a group independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R^1 through R^{13} groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

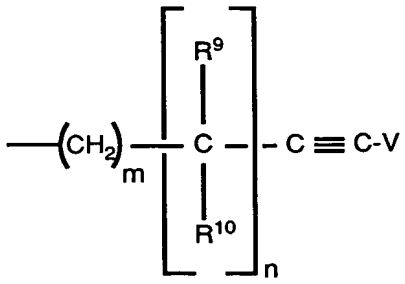
9. The composition of Claim 8 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, iso-butyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'-diethylaminomethyl, N,N'diethylaminoethyl and phenyl; wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R^9 and R^{10} is independently selected from hydrido, alkyl, alkenyl, alkynyl,

cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R^8 is selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl and haloalkenyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkanoyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

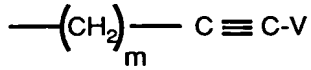
10. The composition of Claim 9 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected from zero through three, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one

through three; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclopropylmethyl, cyclopropylethyl, propylcarbonyl, ethylcarbonyl, methylcarbonyl, phenyl, benzyl, phenylethyl, monochlorophenyl, dichlorophenyl, monofluorophenyl, difluorophenyl, monochlorophenylmethyl, monochlorophenylethyl, dichlorophenylmethyl, dichlorophenylethyl, naphthyl, monofluoronaphthyl, monochloronaphthyl, naphthylmethyl, naphthylethyl, fluoronapthylmethyl and chloronaphthylethyl; or a pharmaceutically-acceptable salt thereof.

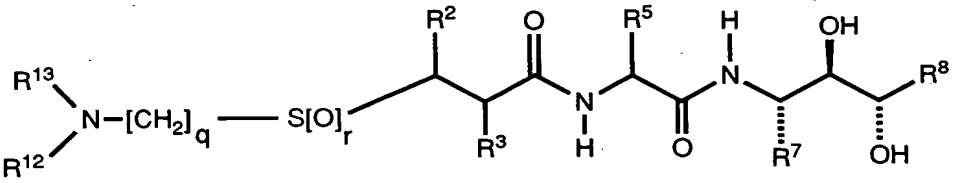
11. The composition of Claim 10 wherein each of R^1 and R^{11} is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from



wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R^{12} and R^{13} is independently

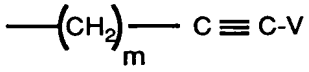
selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

12. The composition of Claim 11 wherein said compound is of Formula II



(II)

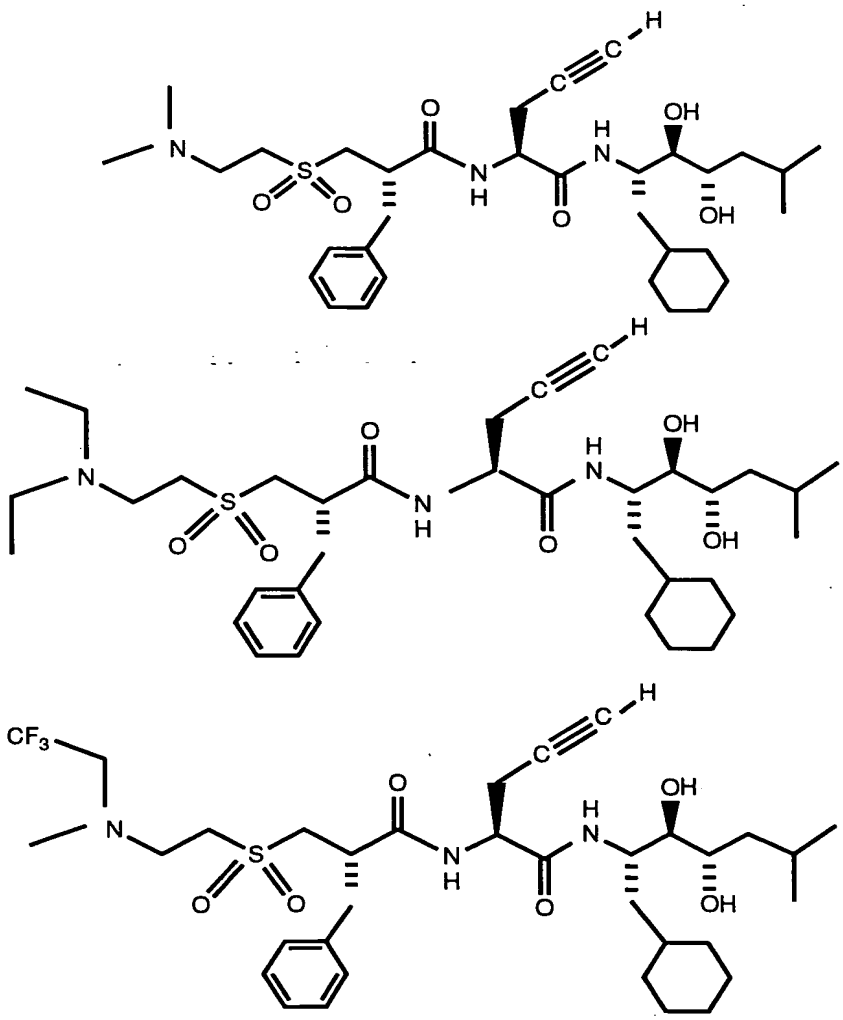
wherein r is zero or two; wherein q is two or three; wherein R^2 is selected from hydrido, methyl, ethyl and phenyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from

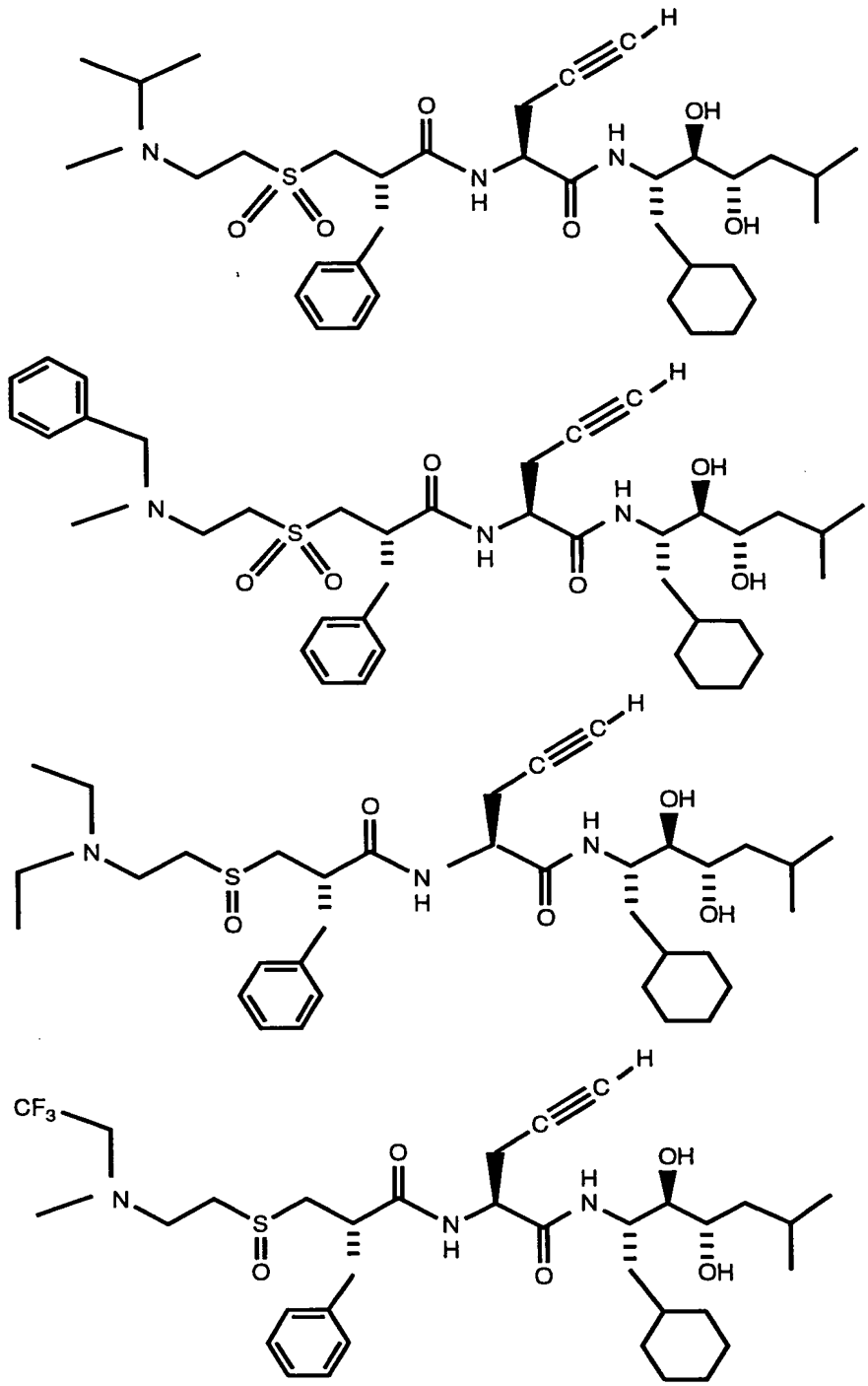


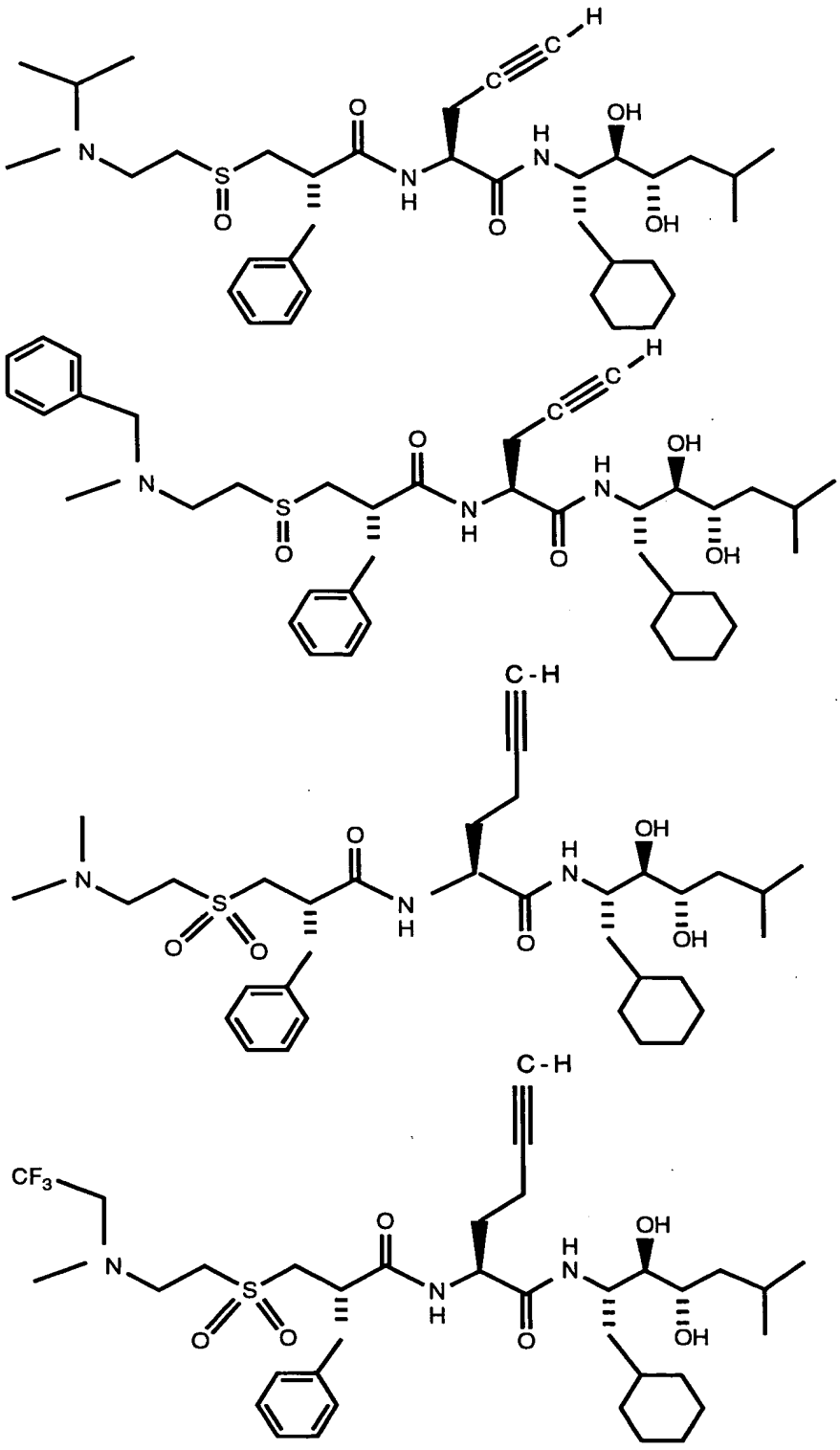
wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is

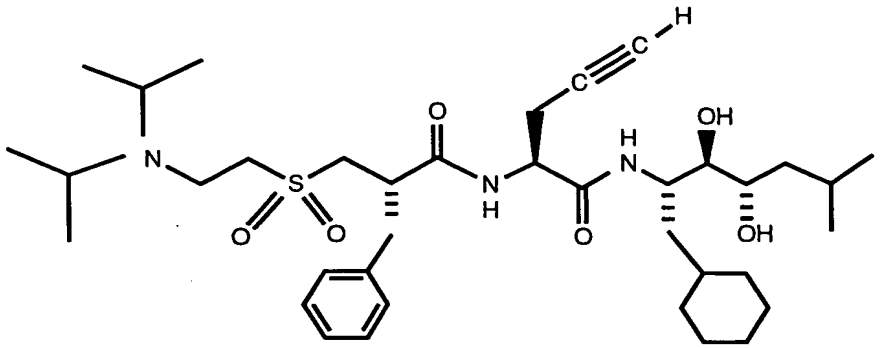
selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R^{12} and R^{13} is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

13. The composition of Claim 12 wherein said renin inhibitor compound is selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

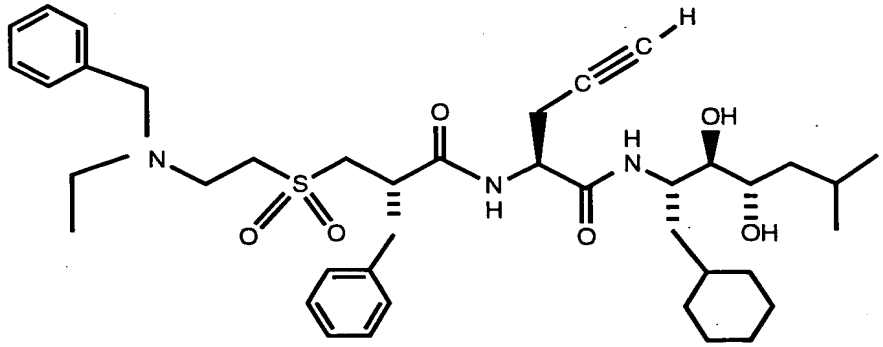






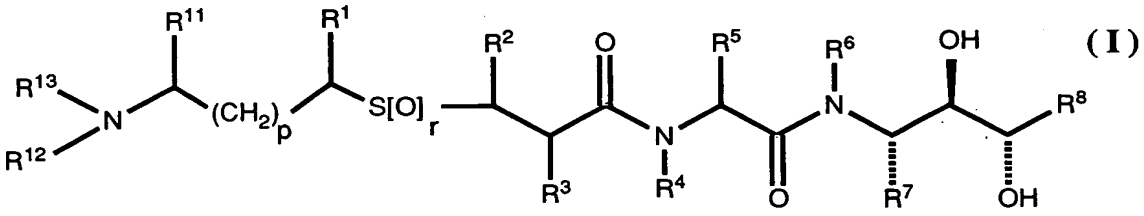


and



14. The composition of Claim 13 wherein said renin-inhibitor compound is N-[1R*-[[[1S,1R*-(cyclohexylmethyl)-2S*,3R*-dihydroxy-5-methylhexyl]amino]carbonyl]-3-butynyl]-αR*-[[[2-dimethylamino)ethyl]sulfonyl]methyl]benzenepropanamide.

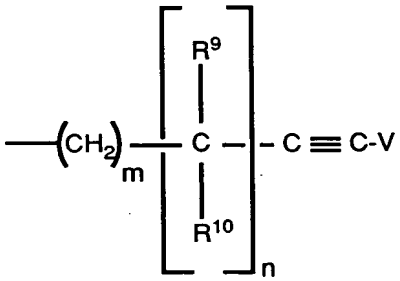
15. A therapeutic method for treating hypertension or glaucoma, said method comprising administering to a hypertensive patient a therapeutically-effective amount of a compound of Formula I:



(I)

wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl;

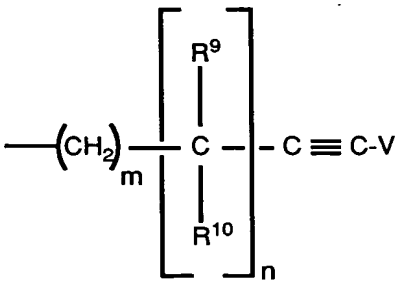
wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is a group selected from hydrido, cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of R^4 and R^6 is a group independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R^9 and R^{10} is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R^8 is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R^{12} and R^{13} is a group independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R^1 through R^{13} groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

16. The method of Claim 15 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, iso-butyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'-diethylaminomethyl, N,N'diethylaminoethyl and phenyl;

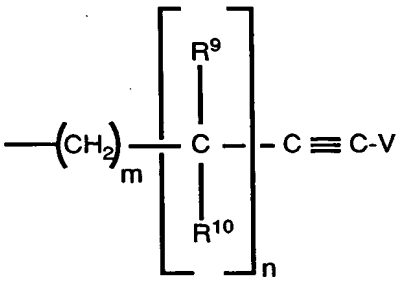
wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R^9 and R^{10} is independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R^8 is selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl and haloalkenyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkanoyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

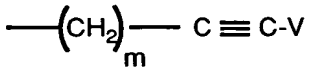
17. The method of Claim 16 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected from zero through three, inclusive; wherein r is

a number selected from zero, one and two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from



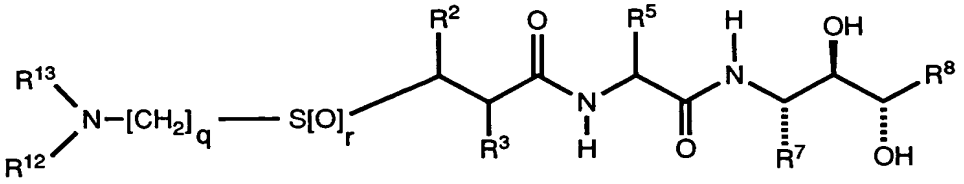
wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one through three; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R^{12} and R^{13} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclopropylmethyl, cyclopropylethyl, propylcarbonyl, ethylcarbonyl, methylcarbonyl, phenyl, benzyl, phenylethyl, monochlorophenyl, dichlorophenyl, monofluorophenyl, difluorophenyl, monochlorophenylmethyl, monochlorophenylethyl, dichlorophenylmethyl, dichlorophenylethyl, naphthyl, monofluoronaphthyl, monochloronaphthyl, naphthylmethyl, naphthylethyl, fluoronapthylmethyl and chloronaphthylethyl; or a pharmaceutically-acceptable salt thereof.

18. The method of Claim 17 wherein each of R^1 and R^{11} is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from



wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R^{12} and R^{13} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

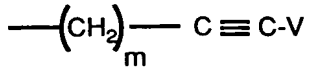
19. The method of Claim 18 wherein said compound is of Formula II



(II)

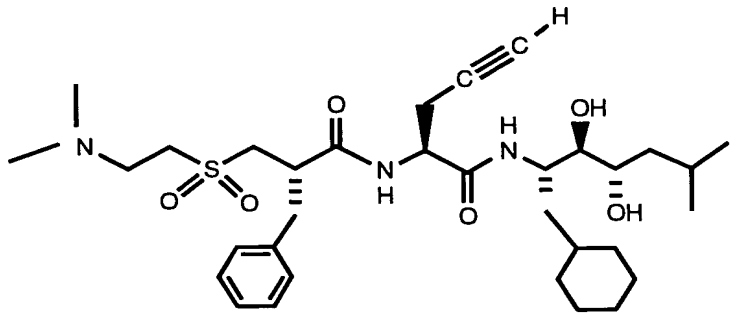
wherein r is zero or two; wherein q is two or three; wherein R^2 is selected from hydrido, methyl, ethyl and phenyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl,

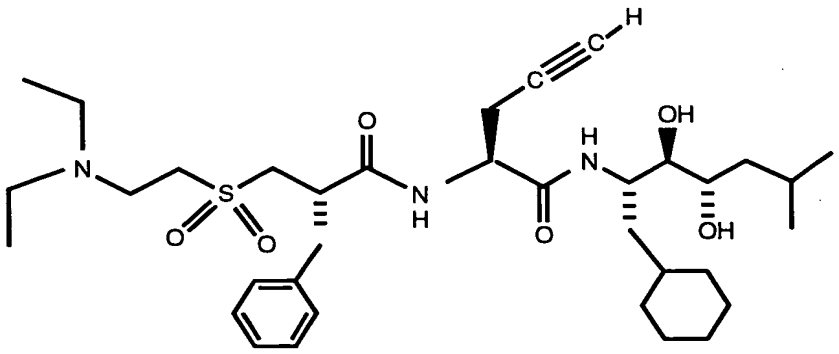
fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from

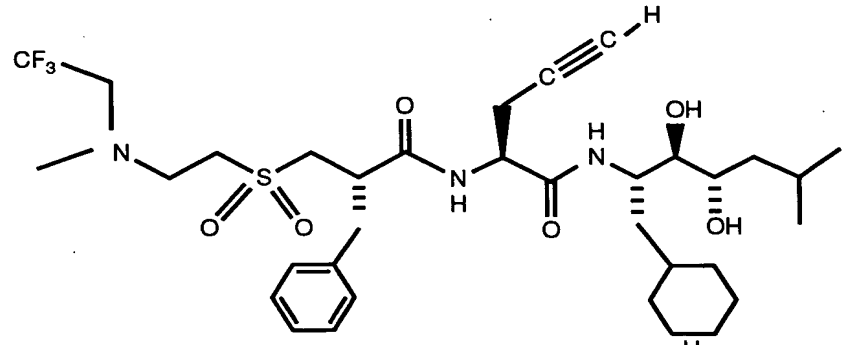


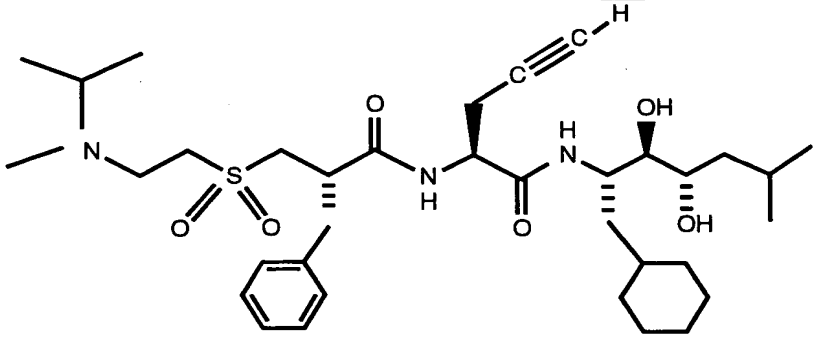
wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R^{12} and R^{13} is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

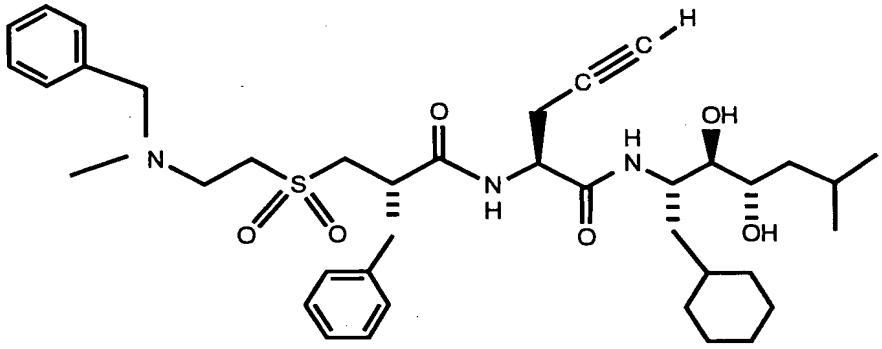
20. The method of Claim 18 wherein said compound is selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

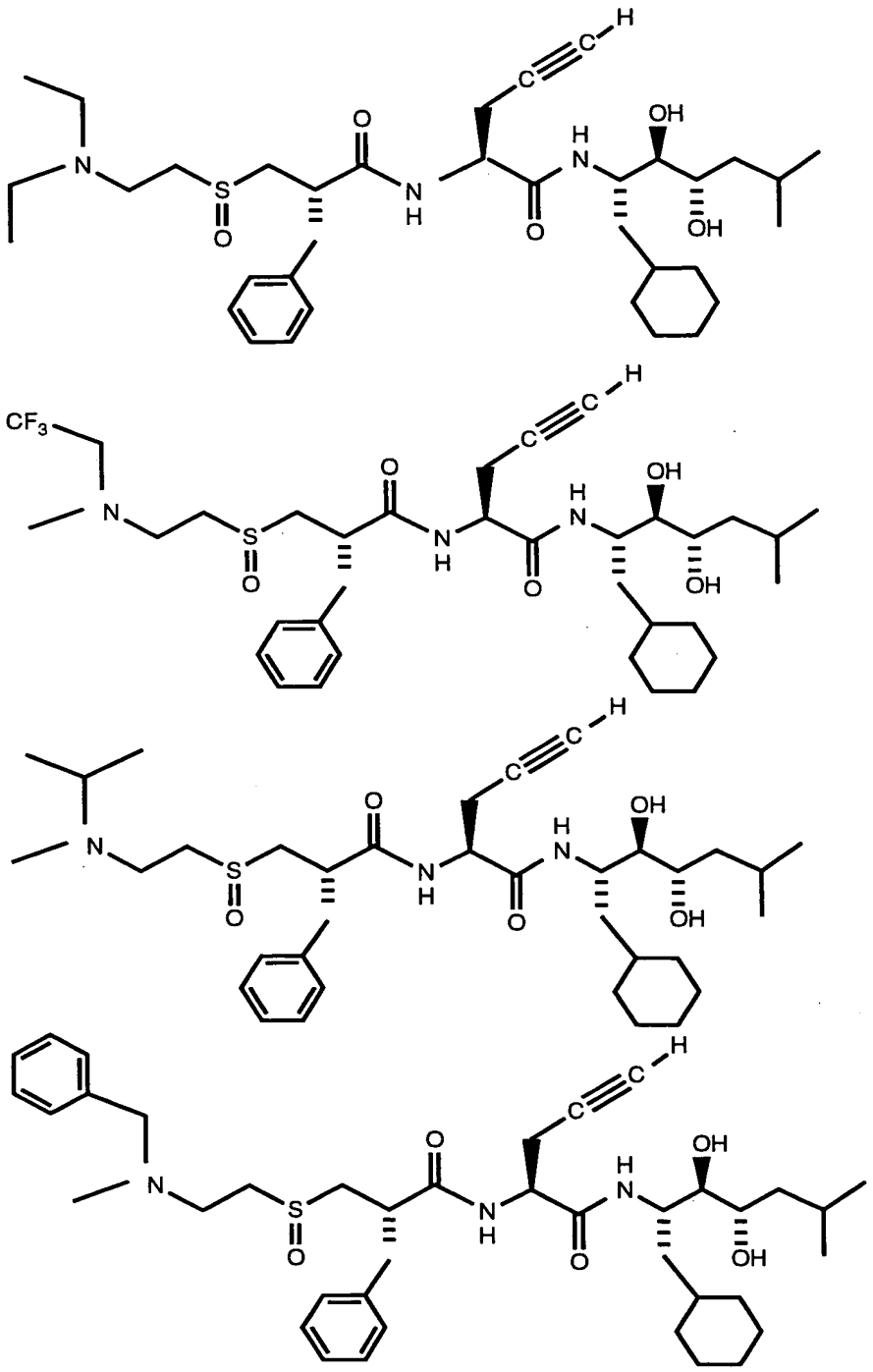


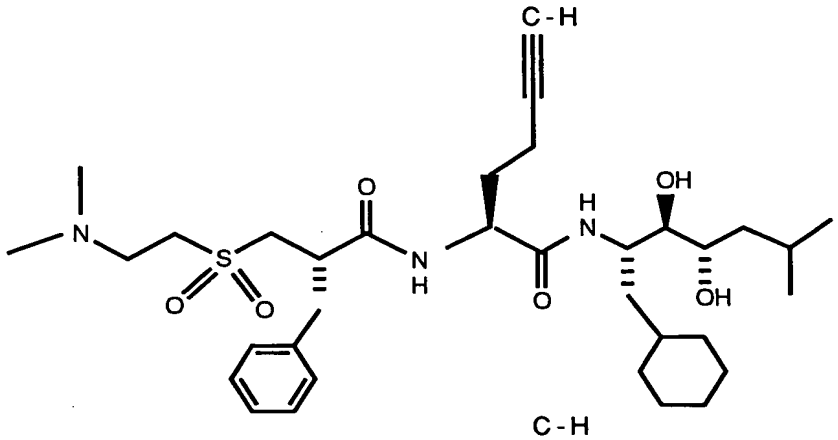


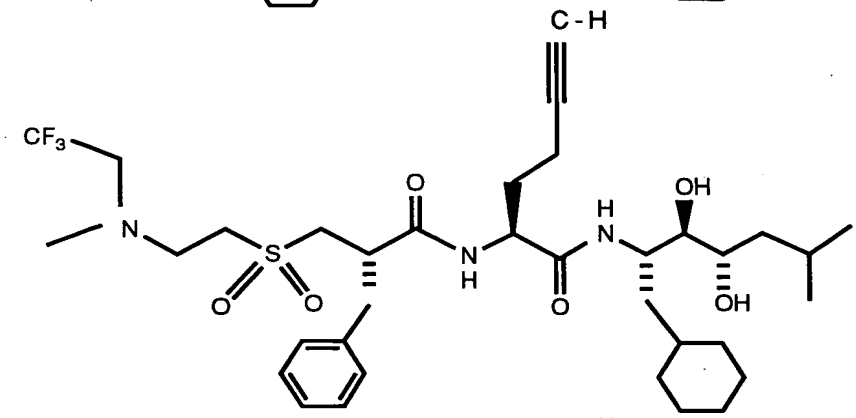


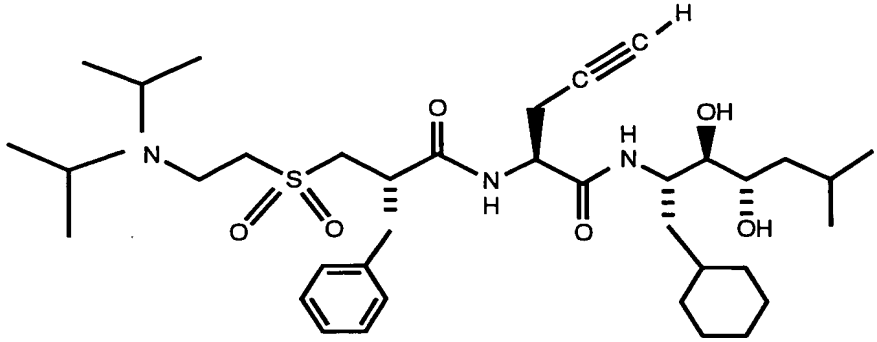




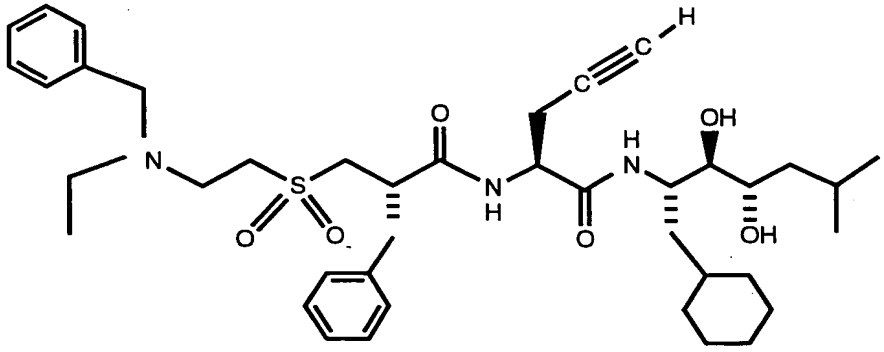








and



21. The method of Claim 20 wherein said compound is N-[1R*-[[[1S,1R*-(cyclohexylmethyl)-2S*,3R*-dihydroxy-5-methylhexyl]amino]carbonyl]-3-butynyl]-αR*-[[[2-(dimethylamino)ethyl]sulfonyl]methyl]benzenepropanamide.

22. The method of Claim 15 for treating hypertension.

23. The method of Claim 15 for treating glaucoma.